

**ENGLISH TRANSLATION  
OF KOREAN PATENT APPLICATION  
NO. 10-2004-0052652**

**KUMKANG INTELLECTUAL PROPERTY LAW OFFICE**

*Re: SN/10/589,328*

[ABSTRACT]

Disclosed herein are a method and kit for diagnosing hereditary diseases CMT1A and HNPP, caused by duplication and deletion in the chromosome 17p11.2-p12 region. In accordance with the present invention, there is provided a method for diagnosing an inherited neuropathy, comprising, running the PCR amplification using microsatellites present in a chromosome 17p11.2-p12 region as markers and DNA typing the resulting PCR amplification products to determine the presence of duplication and deletion in the corresponding chromosomal region, wherein Multiplex PCR amplification is carried out using 6 loci of D17S921, D17S9B, D17S9A, D17S918, D17S122 and D17S4A as markers, and DNA-typing of the resulting PCR amplification products is carried out to determine duplication and deletion in the corresponding chromosomal region. In accordance with the method of the present invention, the diagnosis accuracy of detecting duplication and deletion in the chromosome 17p11.2-p12 region is greater than 99.9%.

[REPRESENTATIVE DRAWING]

Figure 3

[INDEX]

Inherited neuropathy, Chromosome 17, 17p11.2-p12, Duplication, Deletion, CMT1A, HNPP, Diagnosis kits

[DESCRIPTION]

[Invention title]

DIAGNOSIS METHOD AND KITS FOR INHERITED NEUROPATHIES  
CAUSED BY DUPLICATION OR DELETION OF CHROMOSOME 17P11.2-P12  
REGION

[Description of the Drawings]

The above and other objects, features and other advantages of the present invention will be more clearly understood from the following detailed description taken in conjunction with the accompanying drawings, in which:

Fig. 1 shows characteristics of microsatellites distributed in a 1.5 Mb DNA segment of the chromosome 17p11.2-p12 region;

Fig. 2a through 2c show analysis results of PCR products, which were amplified using a diagnostic kit in accordance with the present invention, via an automatic sequencer; and

Fig. 3 shows pedigree charts in which patient pedigrees were analyzed using a diagnostic kit in accordance with the present invention.

[Disclosure]

[Purpose of Invention]

[Technical Field and Background Art]

The present invention relates to a method and kit for diagnosing hereditary diseases CMT1A and HNPP caused by duplication and deletion in the chromosome 17p11.2-p12 region.

Inherited neuropathy (IN) has a relationship with family medical history and is a congenital disease characterized by dysfunction of motor and sensory nervous systems stemming from gene mutations. Among inherited neuropathies, mention may be made of CMT (Charcot-Marie-Tooth disease), HNPP (Hereditary Neuropathy with liability to Pressure Palsies), DSS (Dejerine-Sottas Syndrome) and CH (Congenital Hypomyelination neuropathy), for example. These diseases are caused by genetically heterogeneous factors, and CMT, exhibiting the highest frequency of occurrence among inherited neuropathies, shows an incidence rate of about 1:2,500 in Europeans, and HNPP shows a frequency of occurrence of 16 people per 100,000.

CMT disease is characterized by muscle weakness and atrophy which gradually progresses primarily in peroneal muscle of the lower extremities. This disease may also manifest areflexia, distal sensory loss, pes cavus, and deafness. Onset primarily occurs in the teens, and rarely after the thirties. However, clinical phenotype and onset age of CMT are very heterogeneous and further it is not easy to perform nerve biopsy for definitive diagnosis, thus rendering accurate diagnosis and treatment difficult (Berger P., Young P., Suter U. Neurogenet. 4: 1-15 (2002)).

HNPP is an autosomal dominant-hereditary disease accompanied by repetitive neuralgia and muscle weakness. This disease is also associated with nervous system abnormalities induced by partial thickening of the myelin sheath (tomacula) as revealed by nerve biopsy, and deletion of the chromosome 17p11.2-p12 region is observed in 70%-100% of patients. There are few reported cases of HNPP in Korea, as compared to Western countries, and the reason is considered owing to technical problems in that the presence of tomacula should be confirmed via nerve fiber teasing, in order to make a definitive diagnosis of HNPP disease. However, as it is recently possible to easily

diagnose this disease via genetic testing, it is believed that frequency of such disease cases will increase.

Recently, thanks to studies into the human genome and advanced bioinformatics, studies and research to isolate causative genes responsible for hereditary diseases and to elucidate molecular biological mechanisms thereof are now being actively undertaken throughout the world. A great deal of substantial analysis has been made on hereditary causes of CMT, which is known to be developed by hereditary factors, rather than by environmental factors. As a result, more than 10 genes are known to be involved in pathogenesis of CMT, and most of these mutations have been found to be due to Single Nucleotide Polymorphism (SNP) and duplication/deletion of a short base sequence, except for duplication/deletion of 17p11.2-p12 1.5 Mb. Table 1 below shows inheritance mode and phenotype of important genes responsible for development of CMT known hitherto.

[Table 1]

Genes	Loci	Inheritance mode	Clinical phenotype	No. of mutation found
PMP22	17p.11.2	Autosomal dominant	CMT1, HNPP, DSS, Deafness	59
Cx32X	Xq13.1	X-linked dominant	CMT1X	265
MPZ	1q22	Autosomal dominant Autosomal recessive	CMT1B, CMT2, DSS, Deafness	99
EGR2	10p21.1-p22.1	Autosomal dominant	CMT1(severe), DSS, CH	10
NEFL	8p21	Autosomal dominant Autosomal recessive	CMT1F, CMT2E	24
PRX	10p13.1-p13.2	Autosomal recessive	CMT1, CMT4F, DSS	17

Each CMT patient is caused by a single gene mutation, and pathogenesis thereof very faithfully follows Mendelian inheritance even though there are some differences in severity of symptoms. Since most diseases that currently receive genetic

testing are diseases caused by complicated interaction between polygenic inheritance and various environmental factors, such genetic testing for these diseases simply indicates predisposition to disease pathogenesis (for example, genetic testing of hyperhomocysteinemia-associated MTHFR (5,10-methylenetetrahydrofolate reductase) and breast cancer-inducing BRC-A or BRC-B genes). However, as such, CMT exhibits substantially direct linkage between genetic defects and pathogenesis thereof, as exhibited by Phenylketonuria (PKU) and therefore the results of the genetic testing obtained from such diseases provide a definitive diagnosis, rather than simply at presentation level of predisposition or possibility of disease development, and thus CMT is a genetic disease that is very suitable for utilization of a genetic diagnostic kit.

Therefore, if the etiologies of inherited neuropathies caused by single gene mutation while exhibiting strong hereditary tendency are elucidated via genetic testing, it will be possible to fulfill underlying treatment of diseases in conjunction with more accurate diagnosis. A need for early diagnosis of inherited neuropathies was not raised hitherto due to the absence of relevant treatment methods even though early diagnosis was made for such inherited neuropathies. However, it was recently published in the scientific literature and scientific journals that ascorbic acid and progesterone antagonists are therapeutically effective on CMT1A (Sereda M.W., Horste G.M., Suter U., Uzma N., Nave K.-A. *Nature Genet.* 9: 1533-1537 (2003); Passage E., Norreel J.C., Noack-Fraissignes P., Sanguedolce V., Pizant J., Thirion X., Robaglia-Schlupp A., Pellissier J.F., Fontes M. *Nature Genet.* 10: 396-401 (2004)). Therefore, if methods capable of accurately diagnosing CMT1A duplication/HNPP deletion are developed, inhibition of disease development and patient-tailored treatments may be feasible pursuant to accurate early diagnosis of diseases or diagnosis prior to onset thereof.

The present invention relates to accurate diagnosis of CMT1A and HNPP, pathogenic causes of which are duplication and deletion in the chromosome 17p11.2-p12 region.

Molecular genetic studies into inherited neuropathies were largely led hitherto by European countries, but considerable studies into such diseases are also now being actively undertaken in Japan as well as the USA and European countries since the new millennium. Yoshihara et al. (Yoshihara T., Yamamoto M., Doyu M., Misu K.I., Hattori N., Hasegawa Y., Mokuno K., Mitsuma T., Sobue G. Hum. Mutat. 16: 177-178 (2000)) and Numakura et al. (Numakura C., Lin C., Ikegami T., Guldberg P., Hayasaka K. Human Mutat. 20: 392-398 (2002)) have detected causative mutations in CMT1A duplication, PMP22, MPZ and Cx32 from about 100 Japanese CMT patients. However, there is no accurate statistical data on incidence rate of CMT diseases for Koreans and it is merely believed that situations in Koreans are similar to those of Europeans. The present invention is designed to grasp an accurate definitive diagnosis rate according to a diagnostic method by firstly accurately analyzing hereditary causes of CMT development in Korean CMT patients and their families and comparing the resulting analysis data with foreign data.

CMT which occupied most of inherited neuropathies is broadly classified into CMT1 due to demyelination of the myelin sheath and CMT2 due to an impairment of axon function (*axonopathy*). More than 50% of CMT1 is caused by a 1.5 Mb duplication in the chromosome 17p11.2-p12 region where a PMP22 (peripheral myelin protein 22) gene is located and the resulting disease is called CMT1A disease, taking the highest proportion among overall CMT disease groups. Contrary to CMT1A, HNPP is reported to be due to deletion of the same 1.5-Mb region in 70% to 100% of patients. Therefore, the presence of duplication in the PMP22 gene enables definitive diagnosis

of CMT1A, whereas deletion of the PMP22 gene enables definitive diagnosis of HNPP.

In order to examine CMT1A duplication and HNPP deletion, conventional methods have employed a southern blotting technique using radioisotopes. Recently, via DNA typing of microsatellites present in a 1.5 Mb duplication/deletion region, it became feasible to examine such chromosomal abnormalities without the use of hazardous radioisotopes. However, since markers such as D17S921, D17S955, D17S839 and D17S122, which are largely used in DNA typing, have sequences consisting of (CA)<sub>n</sub> repeat units, PCR of such markers causes severe non-specific DNA amplification due to heavy slippage, thus making it difficult to perform accurate typing of microsatellites (Mersiyanova I.V., Ismailov S.M., Polyakov A.V., Dadali E.L., Fedotov V.P., Nelis E., et al. Human Mutat. 15: 340-347 (2000)). As a result, there were disadvantages associated with necessity of utilizing radioisotopes for accurate diagnosis, and inevitable need of parent's samples for examination of patients.

#### [Technical Problem and Solution]

The present invention relates to diagnosis of CMT1A and HNPP, causes of which are known to be duplication and deletion in the chromosome 17p11.2-p12 region including a PMP22 gene for myelin sheath, respectively.

In the present invention, firstly, samples and clinical data are collected from CMT patient pedigrees and CMT1A duplication and HNPP deletion in patient groups are investigated. Based on these results, relative importance and position of CMT1A duplication and HNPP deletion in onset-diagnosis of overall inherited neuropathies of Koreans, and a rate of definitive diagnosis using the same are provided.

Therefore, the final object of the present invention is to accurately detect



duplication and deletion in the chromosome 17p11.2-p12 region, thereby accurately diagnosing CMT1A and HNPP which are caused by duplication and deletion in the 17p11.2-p12 region.

For the purpose of achieving the above-mentioned objects in accordance with the present invention, new markers including conventional markers are searched and selected as microsatellite markers distributed in the 17p11.2-p12 1.5 Mb region of the short arm of chromosome 17. In particular, as markers such as D17S839 and D17S955, which have been conventionally employed, display 'CA' dinucleotide repeats, PCR amplification thereof leads to heavy slippage. As such, in order to solve such problems and to make accurate diagnosis, the present invention develops and selects new markers having a 3-5 bp repeat sequence, high heterozygosity and various kinds of alleles, among microsatellites present on 17p11.2-p12.

In addition, the present invention constructs primers and constitutes a diagnostic kit, by establishing conditions which enable simultaneous amplification of newly developed markers of interest (Multiplex PCR).

Selected markers are amplified by PCR and the resulting amplification samples are rapidly analyzed by an automatic sequencer, followed by DNA typing using GENESCAN and GENOTYPER programs (Applied Biosystems) to accurately detect duplication and deletion in corresponding regions.

Further, the present invention secures experimental data from comparison and analysis between normal persons (more than 100 individuals) and patient's family groups (more than 50 individuals) utilizing samples and clinical data obtained from CMT patient pedigrees, and constructs standard allele ladders for the respective markers, thus ensuring that accuracy of examination for duplication and deletion in the

corresponding chromosomal regions can be more than 99.9%.

These and other objects and advantages of the present invention will be described hereinafter and will be more fully understood by reference to the following Examples of the present invention.

#### [Construction and Effect of Invention]

In accordance with an aspect of the present invention, the above and other objects can be accomplished by the provision of a method for diagnosing an inherited neuropathy, comprising:

running the Multiplex PCR amplification using microsatellites present in the chromosome 17p11.2-p12 region as markers; and

DNA typing the resulting PCR amplification products to determine the presence of duplication and deletion in the corresponding chromosomal region,

wherein PCR amplification is carried out using 6 loci of D17S921, D17S9B, D17S9A, D17S918, D17S122 and D17S4A as markers, and DNA-typing of the resulting PCR amplification products is then carried out to determine duplication and deletion in the corresponding chromosomal region.

Where duplication and deletion in the 17p11.2-p12 region are determined using the above 6 markers, the accuracy thus obtained is greater than 99.9%. Even with use of some of the above-mentioned markers, for example 3 markers, D17S9B, D17S9A and D17S918 alone, the accuracy of diagnosis is more than 97.5%, which is significantly higher than in conventional diagnostic methods. However, it is preferred to use all 6 markers, in order to reduce the rate of misdiagnosis. Herein, 6 markers may be simultaneously PCR-amplified and used (Hexaplex PCR). If necessary, 6

markers may be used sequentially in a two-step manner involving using 3 markers for diagnosis (Triplex PCR) and using the remaining 3 markers if it is difficult to make definitive diagnosis.

In addition, in accordance with the present invention, there are provided primers having base sequences as set forth in SEQ. ID. NOS: 1-12, in which conditions for simultaneous amplification (Multiplex PCR) were established for the above-mentioned markers.

That is, in one embodiment of the present invention, using a mixture in which primers of SEQ. ID. NOS: 1 and 2; primers of SEQ. ID. NOS: 3 and 4; primers of SEQ. ID. NOS: 5 and 6; primers of SEQ. ID. NOS: 7 and 8; primers of SEQ. ID. NOS: 9 and 10; and primers of SEQ. ID. NOS: 11 and 12 were mixed, the above 6 markers are simultaneously amplified and the resulting amplification products are subjected to DNA typing, thereby determining duplication and deletion in the 17p11.2-p12 region.

In another embodiment of the present invention, the method is carried out in a two-step manner, comprising:

(a) PCR amplification of 3 markers and DNA typing of the resulting PCR amplification products, using a mixture of primers of SEQ. ID. NOS: 3 and 4; primers of SEQ. ID. NOS: 5 and 6; and primers of SEQ. ID. NOS: 7 and 8, thereby firstly determining duplication and deletion in the 17p11.2-p12 region; and

(b) PCR amplification of the remaining 3 markers and DNA typing of the resulting PCR amplification products, using a mixture of primers of SEQ. ID. NOS: 1 and 2; primers of SEQ. ID. NOS: 9 and 10; and primers of SEQ. ID. NOS: 11 and 12, thereby secondly determining duplication and deletion in the 17p11.2-p12 region.

In accordance with another aspect of the present invention, there is provided a kit for diagnosing an inherited neuropathy by determination of duplication and deletion in the chromosome 17p11.2-p12 region, comprising:

a mixture of primers of SEQ. ID. NOS: 1 and 2; primers of SEQ. ID. NOS: 3 and 4; primers of SEQ. ID. NOS: 5 and 6; primers of SEQ. ID. NOS: 7 and 8; primers of SEQ. ID. NOS: 9 and 10; and primers of SEQ. ID. NOS: 11 and 12.

Another form of the diagnostic kit in accordance with the present invention comprises:

a first kit including a mixture of primers of SEQ. ID. NOS: 3 and 4; primers of SEQ. ID. NOS: 5 and 6; and primers of SEQ. ID. NOS: 7 and 8; and

a second kit including a mixture of primers of SEQ. ID. NOS: 1 and 2; primers of SEQ. ID. NOS: 9 and 10; and primers of SEQ. ID. NOS: 11 and 12.

Hereinafter, the present invention will be described in more detail with reference to the following Examples including specific experiments and analyzed cases. These examples are provided only for illustrating the present invention and should not be construed as limiting the scope and spirit of the present invention. Therefore, as will be apparently appreciated by those skilled in the art, various modifications, additions and substitutions are possible, without departing from the scope and spirit of the invention as disclosed in the accompanying claims.

#### **Collection of patient's samples and DNA extraction therefrom**

##### **(1) Collection of samples and clinical data from CMT patient pedigrees**

The present invention has collected and analyzed samples from about 150

pedigrees including Korean CMT patients and their families. Samples of patients and their families were those collected from hospitals for about 10 years from the year 1993, while securing clinical manifestations of patients and past medical history of the patient pedigrees. Based on the thus-obtained samples and clinical data, mutual relationship between clinical features of individual patients and genetic mutations was investigated.

### **(2) Sample collection and DNA extraction**

Blood or saliva, as a sample, was collected from members of pedigrees of patients suffering from inherited neuropathies, and normal persons (control). After blood was pooled in a tube treated with EDTA, genomic DNA was extracted using a DNA purifier. In addition, DNA extraction from hair roots or saliva of subjects was carried out by treating hair or saliva with Proteinase K at a temperature of 55°C for 3 hours, followed by phenol:chloroform extraction.

### **(3) Assessment of FDS (functional disability scale) due to CMT**

In order to determine severity of CMT based on clinical manifestations and the like, the functional handicap was assessed according to a nine-point functional disability scale (FDS) as described by Birouk et al., 1998: 0 = normal; 1 = normal, but with presence of cramps and fatigability; 2 = inability to run; 3 = walking difficult but still possible unaided; 4 = able to walk with a cane; 5 = able to walk with crutches; 6 = able to walk with a walker; 7 = wheelchair bound; 8 = bedridden.

### **Selection of Markers**

In order to exactly detect duplication and deletion in the chromosome 17p11.2-

p12 region, new markers, including markers which have been used in studies hitherto, were searched as microsatellite markers distributed in the 17p11.2-p12 1.5 Mb region of the short arm of chromosome 17. In particular, as markers such as D17S839 and D17S955, which have been conventionally employed, contain 'CA' dinucleotide repeats, PCR amplification of such markers thus leads to heavy slippage. Therefore, the present invention has selected tri- to penta nucleotide markers.

Fig. 1 shows characteristics of microsatellites distributed in a 1.5 Mb DNA segment of the chromosome 17p11.2-p12 region. The present invention has selected markers having high heterozygosity ( $H \geq 0.7$ ) and various kinds of alleles, among various microsatellites present on 17p11.2-p12. As a result, microsatellites of 6 loci including D17S921: (CA)<sub>n</sub>, D17S9B: (TTTC)<sub>n</sub>, D17S9A: (CAATA)<sub>n</sub>, D17S918: (CAG)<sub>n</sub>, D17S122: (CA)<sub>n</sub>, and D17S4A: (ACCT)<sub>n</sub> were finally selected as markers (See Table 2 below).

#### **Primer construction and Kit fabrication**

##### **(1) Establishing conditions for simultaneous amplification (Multiplex PCR)**

For PCR amplification of microsatellite markers, fluorescence-labeled primers were constructed. The primers were designed to have a  $T_m$  value of about 60 to 65 °C similar to those of selected markers. Where there were overlaps in sizes of PCR amplification products between respective markers, primers were labeled with two different fluorescent dyes (FAM and HEX). Table 2 below shows respective primer sequences corresponding to the selected markers.

[Table 2]

Markers	Repeat unit	Dye-labeling	Length	Primer sequences
D17S921	(CA) <sub>n</sub>	FAM	133-149	F: GTGTTGTATTAGGCAGAGTTCTCC (SEQ.ID.NO:1) R: GGCAGTAGATGGTGACTTTATGGC (SEQ.ID.NO:2)
D17S9B	(TTTC) <sub>n</sub>	HEX	95-135	F: TCTCAGTCCTGATTTCTTGATTTTG (SEQ.ID.NO:3) R: CCAGAGCTAACACCACATTCA (SEQ.ID.NO:4)
D17S9A	(CAATA) <sub>n</sub>	HEX	154-204	F: CAACCATCAGTGATTTGATGGTTTAC (SEQ.ID.NO:5) R: GAGTTGTCACTAGAACCTGTTC (SEQ.ID.NO:6)
D17S918	(CAG) <sub>n</sub>	HEX	235-250	F: TCCTGTAATCTGTCCCAAACGTC (SEQ.ID.NO:7) R: TTCCTCACACAACCTATTGATAGTC (SEQ.ID.NO:8)
D17S122	(CA) <sub>n</sub>	FAM	206-228	F: AGAGCTGTTCTGCTGAAGTCACTC (SEQ.ID.NO:9) R: GGCAGGCCAGACAGACCAGGCTC (SEQ.ID.NO:10)
D17S4A	(ACCT) <sub>n</sub>	FAM	160-180	F: CTGTGGAGGAAAGAAAACACTGCC (SEQ.ID.NO:11) R: GCACTAAAGTAGCTTGTAACCTCG (SEQ.ID.NO:12)

## (2) Construction of PCR Kit

A PCR kit in accordance with the present invention is constructed in the form of a Hexaplex PCR kit that performs simultaneous amplification of 6 markers as listed in Table 2, or a triplex PCR kit that performs simultaneous amplification of 3 respective markers. In addition, experimental data from comparison and analysis between normal persons (n = 100) and patient's family groups (n = 50), and standard allele ladders for respective markers were prepared such that the accuracy of examination was greater than 99.9%. Preparation of the PCR kits is as follows.

### Preparation example 1

In order to simultaneously amplify all 6 locus markers, i.e., D17S921, D17S9B, D17S9A, D17S918, D17S122 and D17S4A, a Hexaplex PCR kit was constructed including a mixture of 6 pairs of primers in differential concentrations, wherein the

respective primers were represented by SEQ. ID. NOS: 1 through 12.

### **Preparation example 2**

In this example, a PCR kit was constructed to include Kit A and Kit B as triplex PCR kits, respectively. Kit A includes a mixture of 3 pairs of primers (SEQ. ID. NOS: 3 through 8) labeled with HEX in differential concentrations, in order to amplify markers of 3 loci, i.e., D17S9A, D17S9B and D17S918. Whereas, Kit B includes a mixture of 3 pairs of primers (SEQ. ID. NOS: 1 and 2, 9 through 12) labeled with FAM in differential concentrations, in order to amplify markers of 3 loci, i.e., D17S921, D17S122 and D17S4A. Since respective 3 markers constituting Kits A and B were labeled with the same fluorescent materials, genotypes thereof are confirmed on single chromatogram upon analyzing genotypes of such markers.

### **Genotyping and diagnosis utilizing diagnostic kit**

After PCR was carried out using a diagnostic kit in accordance with the present invention, an isolation system using a single-lane was applied to each sample by means of an automatic sequencer, followed by DNA typing using GENESCAN and GENOTYPER programs.

### **(1) Application of DNA diagnostic kit for diagnosing CMT1A and HNPP to patient pedigrees**

Using the kit in accordance with the present invention, genetic testing was carried out on 3 families of patients affected with true inherited neuropathies (FC87, FC91, and HN42).



### **① Clinical analysis of analyzed patient pedigrees**

FC87 : 'Mother' and 'offspring (son)' are CMT1 patients; and 'father' is normal.

FC91 : 'Mother', 'offspring 1 (son)' and 'offspring 2 (daughter)' are CMT1 patients; and 'father' is normal.

HN42 : 'Father' and 'offspring' are HNPP patients; and 'mother' is normal.

### **② Diagnostic kit**

Markers obtained from the above-mentioned FC87, FC91 and HN42 patient pedigrees were amplified using a Hexaplex PCR kit including a mixture of 6-paired primers (SEQ. ID. NOS: 1 to 12) and a standard allele ladder, as prepared in Preparation Example 1.

### **③ Conditions of PCR amplification**

96 °C (3min)

94 °C (1min)-60 °C (1min)-70 °C (1min): 32 cycles

60 °C (50min)

### **④ Analysis**

The resulting PCR amplification products were analyzed using an automatic sequencer (ABI 3100) and genotyping thereof was carried out using GENESCAN and GENOTYPER programs (Applied Biosystems, USA).

### **⑤ Results**

Results are shown in Fig. 2, Fig. 3 and Table 3 below.

Fig. 2 shows analysis results of PCR products amplified by the diagnostic kit in accordance with the present invention, using the automatic sequencer. Fig. 2a shows analysis results of PCR products for CMT1 patient's family (FC 87), thus representing no duplication and deletion in 17p11.2-p12. Fig. 2b shows analysis results of PCR products for CMT1 patient's family (FC 91), thus representing 'duplication' in the corresponding chromosomal region. Fig. 2c shows analysis results of PCR products for HNPP patient's family (HN42), thus representing 'deletion' in the corresponding chromosomal region.

Table 3 below summarizes data of microsatellite genotyping obtained from results of Fig. 2.

[Table 3]

Pedigree No.	Relation	Sex	Phenotype	Microsatellite markers						Remark
				D17S921	D17S98	D17S9A	D17S918	D17S122	D17S4A	
FC87 (CMT1)	Father	Male	Normal	6, 7	11, 11	11, 14	8, 8	3, 11	16, 19	
	Mother	Female	Affected	3, 7	12, 15	13, 13	8, 12	3, 11	14, 2, 18	
	Son	Male	Affected	7, 7	11, 12	13, 14	8, 8	3, 3	18, 19	
FC91 (CMT1)	Father	Male	Normal	3, 6	13, 14	13, 13	12, 12	12, 14	14, 2, 15, 2	
	Mother	Female	Affected	6-7, 7	12-12, 15	13-15, 12	8-12, 8	3-13, 12	14, 2-18, 17	Duplication
	Son	Male	Affected	6-7, 3	12-12, 13	13-15, 13	8-12, 12	3-13, 12	14, 2-18, 15, 2	Duplication
	Daughter	Female	Affected	6-7, 3	12-12, 13	13-15, 13	8-12, 12	3-13, 12	14, 2-18, 15, 2	Duplication
NH42 (HNPP)	Father	Male	Affected	4, (-)	9, (-)	14, (-)	12, (-)	12, (-)	16, (-)	Deletion
	Mother	Female	Normal	3, 7	11, 11	11, 13	8, 12	11, 12	14, 2, 16	
	Son	Male	Affected	7, (-)	11, (-)	13, (-)	8, (-)	12, (-)	16, (-)	Deletion

Based on the above analysis results, pedigree charts of 3 families are shown in Fig. 3 (□, ○: Normal; ■, ●: Affected).

The analysis results are summarized according to respective pedigrees as follows.

**(a) FC87 pedigree**

When the kit of the present invention was applied, duplication or deletion in the chromosome 17p11.2-p12 region was not observed in this patient family. Therefore, it was determined that pathogenesis of this family was due to mutation in other genes, other than duplication/deletion in the 17p11.2-p12 region.

**(b) FC91 pedigree**

From genetic testing via application of the kit in accordance with the present invention, duplication in the 17p11.2-p12 region was observed in 'mother', 'offspring 1' and 'offspring 2' of the FC91 patient family. Therefore, 'mother', 'offspring 1' and 'offspring 2' were diagnosed as CMT1A patients. In addition, analysis results showed that there was distinctive consistency in "chromosomal duplication-patient relationship," inherited from the patient 'mother' to two offsprings (patients). As shown in Fig. 3, shaded parts represent duplicated chromosome regions that were found in mother, offspring 1 and offspring 2.

**(c) HN42 pedigree**

From genetic testing via application of the kit in accordance with the present invention, deletion in the 17p11.2-p12 region was observed in 'father' and 'offspring' of the HN42 patient family, while duplication or deletion in the corresponding chromosome region was not observed in 'mother'. That is, it was confirmed that

chromosomal deletion of 'father' was inherited by offspring, thereby rendering offspring to be HNPP patients.

**(2) Investigation into CMT1A duplication and HNPP deletion for Korean patient groups**

Duplication in 17p11.2-p12 of chromosome 17 was investigated by genotyping 6 microsatellite markers in 28 families of Korean CMT1 patients. As a result, segmental duplication of chromosome 17 was observed in 15 families, thus representing a duplication rate of 53.6%. The duplication rate investigated in the present invention shows the results similar to 57.6% (Mostacciuolo et al. Human Mutat. 18: 32-41 (2001) in Italy, and 53.7% (Mersiyanova et al. Human Mut. 15: 340-347 (2000)) in Russia. In addition, segmental deletion of chromosome 17, which is responsible for HNPP, was observed in 19 families out of the total 24 families, thus representing a deletion rate of 79%.

**[Industrial Applicability]**

The present invention enables a diagnosis accuracy of greater than 99.9% in detection of duplication and deletion in the chromosome 17p11.2-p12 region. In addition, in accordance with the present invention, it is possible to make definitive diagnosis of exact hereditary causes for more than 50% of CMT patients and more than 70% of HNPP patients upon first diagnosis using the diagnostic kit in accordance with the present invention, when considering an incidence rate of inherited neuropathies, pathogenic causes of which are duplication and deletion in the chromosome 17p11.2-p12 region. Further, co-application of the diagnostic kit in

accordance with the present invention, in conjunction with a diagnostic kit for diagnosing an inherited neuropathy caused by other gene mutations except for duplication and deletion in the chromosome 17p11.2-p12 region, which is another invention of the present inventors, enables a detection rate of more than 90% of total hereditary causes. In addition, the present invention provides an advantage such as no use of radioisotopes by excluding conventional southern blotting. Therefore, the present invention enables exact early diagnosis of inherited neuropathies caused by single gene mutation while exhibiting strong hereditary predisposition via genetic testing, early application of recently-developing treatment methods pursuant to exact diagnosis, and further patient-tailored therapy corresponding to exact etiological causes.

**[CLAIMS]**

[Claim 1] deleted

[Claim 2] deleted

[Claim 3] deleted

[Claim 4] A kit for diagnosing an inherited neuropathy by determination of duplication and deletion in the chromosome 17p11.2-p12 region, comprising:

a mixture of primers of SEQ. ID. NOS: 1 and 2; primers of SEQ. ID. NOS: 3 and 4; primers of SEQ. ID. NOS: 5 and 6; primers of SEQ. ID. NOS: 7 and 8; primers of SEQ. ID. NOS: 9 and 10; and primers of SEQ. ID. NOS: 11 and 12.

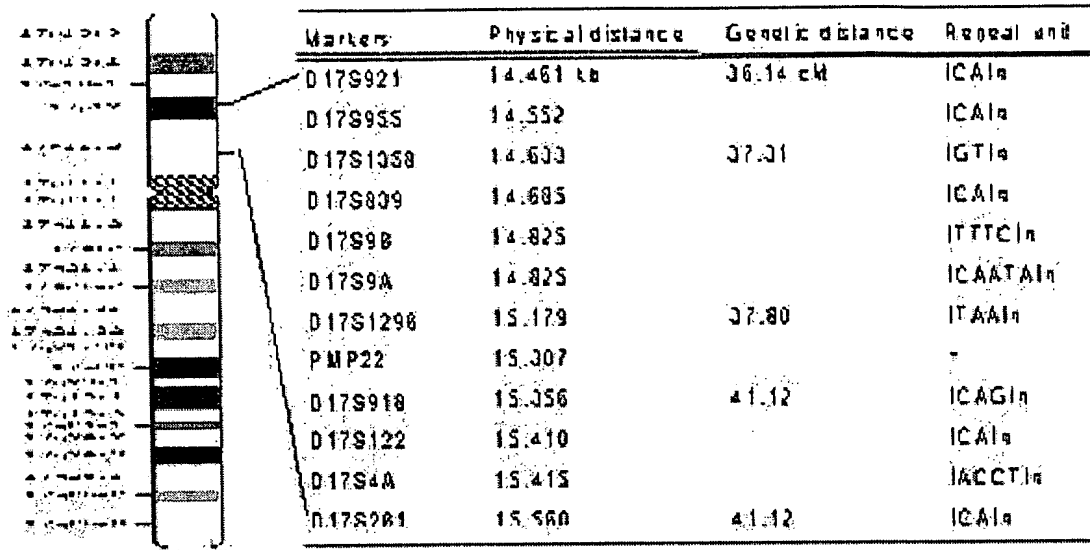
[Claim 5] A kit for diagnosing an inherited neuropathy by determination of duplication and deletion in a chromosome 17p11.2-p12 region, comprising:

a first kit including a mixture of primers of SEQ. ID. NOS: 3 and 4; primers of SEQ. ID. NOS: 5 and 6; and primers of SEQ. ID. NOS: 7 and 8; and

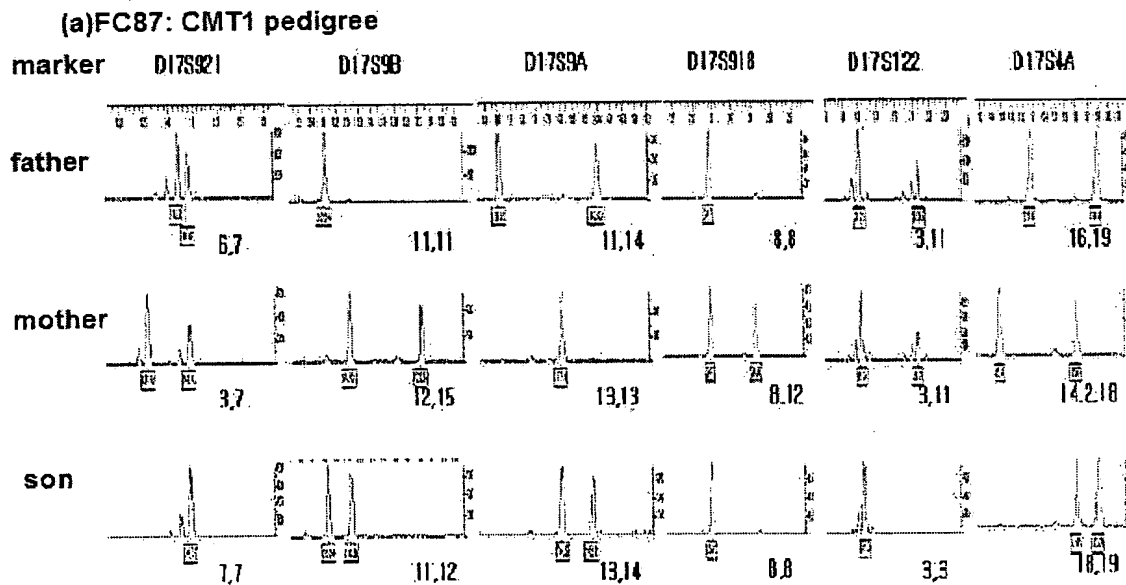
a second kit including a mixture of primers of SEQ. ID. NOS: 1 and 2; primers of SEQ. ID. NOS: 9 and 10; and primers of SEQ. ID. NOS: 11 and 12.

# [DRAWINGS]

[Figure 1]

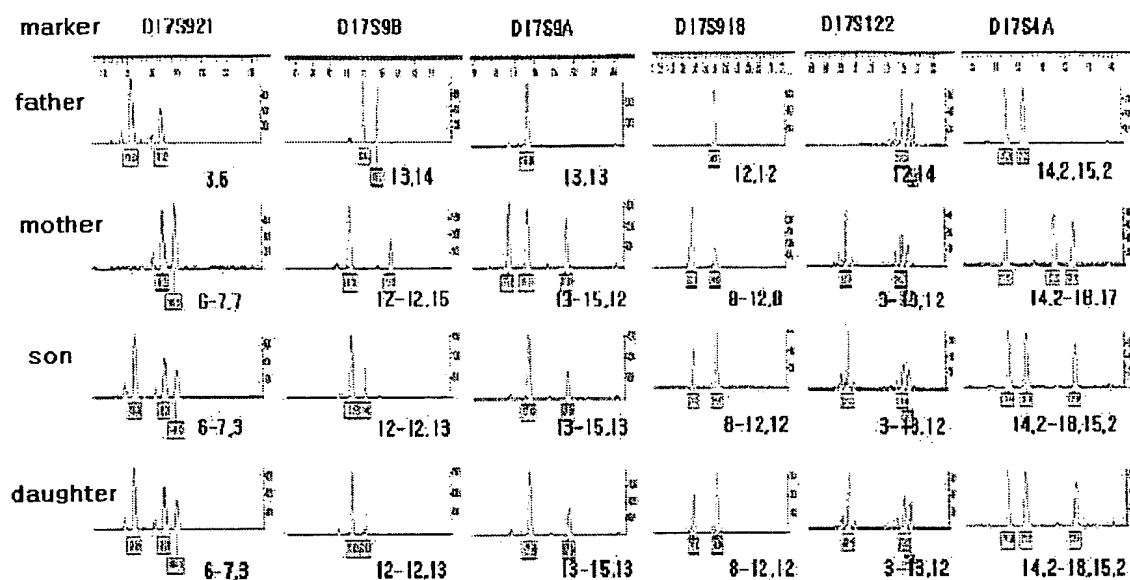


[Figure 2a]



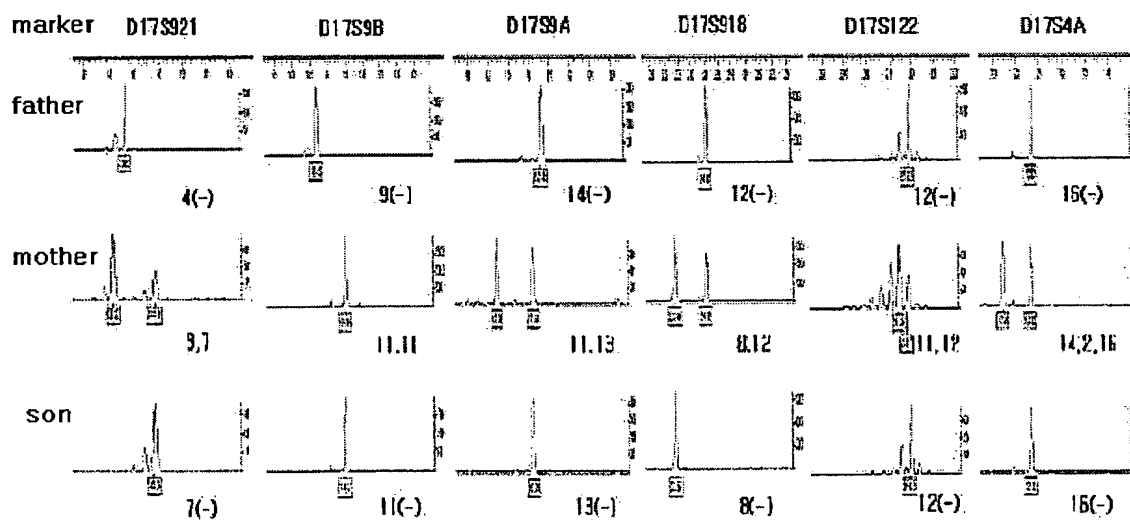
[Figure 2b]

(b)FC91: CMT1 pedigree



[Figure 2c]

(c)HN42: HNPP pedigree





[Figure 3]

(a)FC87 CMT1 pedigree (b)FC91 CMT1 pedigree (c)HN42 HNPP pedigree

